

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
18 April 2002 (18.04.2002)

PCT

(10) International Publication Number
WO 02/31085 A2

(51) International Patent Classification⁷: **C10G 21/04**

(74) Agents: GILSON, David, Grant et al.; Spoor and Fisher,
P.O. Box 41312, 2024 Craighall (ZA).

(21) International Application Number: PCT/IB01/01874

(22) International Filing Date: 9 October 2001 (09.10.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:

2000/5508 9 October 2000 (09.10.2000) ZA
60/239,060 9 October 2000 (09.10.2000) US

(71) Applicant (for all designated States except US): SASOL
TECHNOLOGY (PTY) LTD. [ZA/ZA]; 1 Klasie
Havenga Road, 1947 Sasolburg (ZA).

(72) Inventors; and

(75) Inventors/Applicants (for US only): DE WET, Johan,
Peter [ZA/ZA]; 65 Beethoven Street, 1911 Vanderbijlpark
(ZA). SCHOLTZ, Jacob, Johannes [ZA/ZA]; 6 Oost-
huizen Street, 1947 Sasolburg (ZA).

(81) Designated States (national): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,
MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK,
SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA,
ZW.

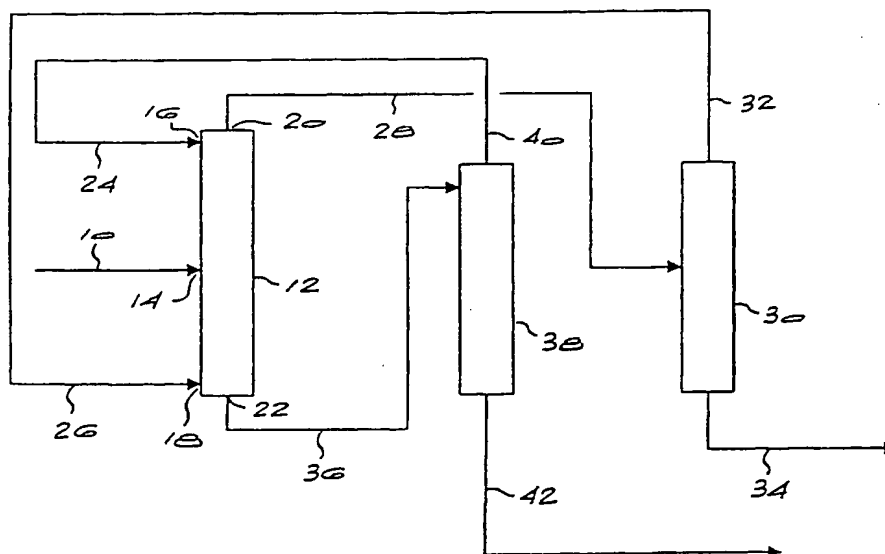
(84) Designated States (regional): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF,
CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD,
TG).

Published:

— without international search report and to be republished
upon receipt of that report

[Continued on next page]

(54) Title: SEPARATION OF OXYGENATES FROM A HYDROCARBON STREAM



(57) Abstract: This invention relates to a method for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream containing a high proportion of olefins, paraffins and oxygenates (mainly alcohols). Typically, the hydrocarbon stream is obtained from a Fischer-Tropsch process. The organic counter-solvent has a boiling point which is less than the boiling point of the most volatile alcohol in the hydrocarbon stream. A raffinate from the liquid-liquid extractor is passed to a distillation column. A bottoms product from the distillation column comprises olefins and paraffins, and the overhead product comprising solvents is recycled. An extract from the liquid-liquid extractor is sent to a stripping column, where a bottoms product containing pure alcohol is obtained. The overhead product containing counter-solvent is recycled.

WO 02/31085 A2



For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

-1-

SEPARATION OF OXYGENATES FROM A HYDROCARBON STREAM

BACKGROUND OF THE INVENTION

THIS invention relates to a process for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream.

The reaction of synthesis gas in a Fischer-Tropsch reactor at elevated temperature and pressure over an Fe or Co catalyst produces a range of hydrocarbons including paraffins, olefins and alcohols, with carbon chain length varying from 1 to greater than 100. The Fischer-Tropsch reaction can occur at a temperature in excess of 300°C. Generally, the operating temperature is in the range of 200°C to 260°C utilising fixed bed or slurry phase reactors. The use of a Fe/Mn/Zn catalyst operated at a pressure of 30-60 bar has been shown to display high selectivity to olefins and oxygenates (mainly alcohols) with chain lengths of from 2 to greater than 30.

CONFIRMATION COPY

-2-

In a prior art processes for the extraction of alpha olefins from Fischer-Tropsch product streams, the concentration of the desired alpha olefin is increased in a first distillation step, followed by etherification, extractive distillation, adsorption and/or superfractionation steps. South African patent no. 98/4676 teaches an improvement in the process for the recovery of alpha-olefins from a mixed hydrocarbon stream derived from the Fischer-Tropsch process. The process described in this prior art involves three distinct steps:

- 1) Coarse separation of hydrocarbon products into a low boiling and a high boiling fraction
- 2) Distillative fine separation of lower and higher boiling fractions
- 3) Etherification of tertiary olefins following the fine separation. In these processes, tertiary olefins were considered as a problematic component that cannot be separated from the desired alpha olefin readily by conventional distillation in order to produce a product that is of polymer grade.

A known commercial process of removing oxygenates from a hydrocarbon stream includes the hydrogenation of a C₁₀ to C₁₃ cut of a hydrocarbon stream containing olefins, paraffins and oxygenates. The hydrogenation step removes oxygenates, but also hydrogenates olefins to paraffins, which is undesirable.

US patent no. 4,686,317 discloses a process for removing oxygenated impurities from a light (C₂ to C₄) hydrocarbon stream which includes extracting the oxygenates with a heavy organic polar solvent, water scrubbing the extracted hydrocarbons to recover the dissolved solvent, and combining the solvent phase from the extraction and water phase from the scrubber and distilling to recover the solvent. This process is not concerned with the recovery of oxygenates from the extract.

SUMMARY OF THE INVENTION

According to the invention there is provided a method for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream including hydrocarbons and oxygenates, the method including contacting the hydrocarbon stream under conditions of liquid-liquid extraction with a polar solvent and a non-polar organic counter-solvent, wherein the organic counter-solvent comprises olefins and/or paraffins.

Preferably, the organic counter-solvent is selected to have a boiling point which is less than the boiling point of the most volatile alcohol in the hydrocarbon stream.

Advantageously, the hydrocarbon stream is passed through a distillation column prior to introduction to the liquid-liquid extraction, to remove C_1 to C_7 hydrocarbons and C_1 to C_3 alcohols from the stream. In this case, a suitable organic counter-solvent comprises olefins and/or paraffins in the C_8 range.

Advantageously, C_{21} - plus, more preferably the C_{19} - plus hydrocarbons, are also removed from the hydrocarbon stream by distillation prior to the liquid-liquid extraction so that the stream includes C_4 to C_{20} , preferably C_4 to C_{18} hydrocarbons.

The polar solvent for a C_4 to C_{20} hydrocarbon stream is typically a light polar solvent which advantageously comprises a mixture of water and an organic liquid such as n-propanol. The preferred light polar solvent is however a mixture of water and acetonitrile. Typically, the water comprises no more than the azeotropic composition of water and the organic liquid.

-4-

The liquid-liquid extraction may take place in a liquid-liquid extractor apparatus such as a vertical liquid-liquid extraction column or a mixer-settler type extraction unit.

Raffinate from the liquid-liquid extractor may be introduced into a distillation column and a mixture of paraffins and olefins, with a low concentration of oxygenates, may be recovered as a bottoms product from the distillation column.

The overhead product from the distillation column, which includes counter-solvent and a small amount of solvent, is conveniently recycled to the lower counter-solvent inlet of the extraction column.

Extract from the liquid-liquid extractor may be sent to a stripping column, where the solvent and a small amount of counter-solvent are removed as an overhead product, and then recycled to the upper solvent inlet of the extraction column, and the bottoms product from the stripping column is an oxygenate stream containing low concentrations of non-polar material.

According to a preferred embodiment of the invention there is provided a method for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream comprising C_4 to over C_{20} hydrocarbons and oxygenates, the method including the steps of:

introducing the hydrocarbon stream into a vertical liquid-liquid extraction column, the liquid extraction column including an upper solvent inlet, a lower counter-solvent inlet, an upper raffinate outlet and lower extract outlet;

introducing a polar solvent described above into the extraction column via the upper solvent inlet; and

-5-

introducing a non-polar counter-solvent described above into the extraction column via the lower counter-solvent inlet.

Advantageously, the hydrocarbon stream is added to the liquid-liquid extraction column, along the liquid-liquid extraction column.

BRIEF DESCRIPTION OF THE DRAWING

The drawing is a flow diagram of a process according to the invention.

DESCRIPTION OF EMBODIMENTS

This invention relates to a method for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream comprising C_4 to over C_{20} hydrocarbons under conditions of liquid-liquid extraction with a polar solvent and a non-polar organic counter-solvent. Although the use of a vertical liquid-liquid extraction column is described, any type of liquid-liquid extraction unit, such as a mixer-settler unit, may be used.

Referring to the diagram, a liquid hydrocarbon stream 10 (hereinafter referred to as the "feed") containing C_4 to over C_{20} , preferably C_8 to C_{20} , typically C_8 to C_{18} , olefins and paraffins, and impurities such as oxygenates (which may include C_4 to C_{16} alcohols, ethers, aldehydes, ketones, acids and mixtures thereof) is fed into a vertical liquid-liquid extraction column 12 via a feed inlet 14 along the column. The extraction column 12 includes an upper solvent inlet 16, a lower counter solvent inlet 18, an upper raffinate outlet 20 and a lower extract outlet 22.

-6-

The feed may be obtained from a Fischer-Tropsch reaction. In a preferred embodiment, the hydrocarbon stream is the product of a Fischer-Tropsch reaction in a slurry bed reactor using an Fe/Mn/Zn catalyst operated at a pressure of 30-65 bar and at a temperature range of 200°C to 260°C. The reactor conditions are selected to provide a high proportion of olefins and paraffins, and alcohols. Typically, the condensate comprises olefins, paraffins and up to 55% by mass alcohols. The hydrocarbon stream is passed through a distillation column prior to introduction to the liquid extraction column 12, to remove C₁ to C₇ hydrocarbons and C₁ to C₃ alcohols from the stream. An additional distillation column can also be used to remove the heavier hydrocarbons, i.e. C₂₁ - plus hydrocarbons, if desired. The removal of the light (C₁ to C₃) alcohols is important, otherwise it will be difficult to remove them from the extract 36 described below.

A polar solvent 24, which in this embodiment of the invention is a light polar solvent mixture of water and acetonitrile, is introduced to the column 12 via the upper inlet 16. By light polar solvent, it is meant that the solvent should be immiscible with the hydrocarbons and have a boiling point lower than that of the oxygenates and the hydrocarbons. Another light solvent that is suitable for hydrocarbon stream containing a wide range of hydrocarbons, i.e. from C₄ to C₂₀, is a water/n-propanol mixture. A light polar solvent which comprises a mixture of water and acetonitrile, with no more than 19%, by mass, water is however preferred. Water forms a light-boiling azeotrope with acetonitrile. In the present conditions, the water-acetonitrile azeotrope will be the lightest boiling item in the extract. This is advantageous as it aids solvent recovery in the stripper 38 described below. If the water content of the acetonitrile exceeds that of the water acetonitrile azeotrope concentration, there will be excess water after the water-acetonitrile azeotrope has been boiled off. If there are any other components in the extract that may form light-boiling azeotropes with water, it will do so in the presence of the excess water. Such azeotrope(s) may also be boiled to the overheads, and may leave with the

-7-

solvent. The danger exists that such components may then accumulate in the solvent loop, and this in turn may be detrimental to the efficiency of the separation.

The ratio of solvent to feed must be sufficient to exceed the solubility of the solvent in the hydrocarbons to form two distinct liquid phases. Usually, the solvent to feed ratio will be from 2:1 to 6:1. A low solvent to feed ratio is preferred as less solvent needs to be recovered in the solvent stripper 38 mentioned below. This saves energy by boiling off less solvent as an overhead product.

An organic counter-solvent 26, is introduced to the extraction column 12 via the lower inlet 18. It is important that the counter solvent has a boiling point which is low enough for it to be separated from the most volatile alcohols present in the stream, in the stripper 38 which is described below. In the present case, where the C₁ to C₃ alcohols have been removed, the boiling point of the counter-solvent must be low enough to be separated from C₄ alcohols (n-butanol). A preferred counter-solvent comprises an olefin or paraffin, typically a mixture of olefins and paraffins mixture, in the C₈ range. A typical counter-solvent is octene.

The ratio of feed to the counter-solvent should be approximately 2:1. A preferred solvent to counter-solvent ratio is from 1:2 to 1:6, typically 1:4.

A raffinate 28 leaving the upper exit 20 of the extraction column 12 consists of non-polar hydrocarbons, counter-solvent and a small amount of solvent. The raffinate 28 is introduced into a distillation column 30. Counter-solvent and a small amount of solvent is recovered from the raffinate as an overhead product of the distillation column and is recycled to the lower inlet 18 of the extraction column 12. A bottoms product 34 from the distillation column 30 yields a mixture of olefins and paraffins, with low a concentration of oxygenates.

-8-

An extract 36 from the lower outlet 22 of the extraction column 12 is sent to a stripper 38. An overhead product 40 from the stripper 38, which contains solvent and small amount of counter solvent is recycled through the upper inlet 16 of the extraction column 12 (it may be necessary to replenish this solvent with a make-up stream to counter losses). A bottoms product 42 from the stripper 38 has a high concentration of oxygenates, and low concentration of non-polar material. As mentioned above, any counter-solvent in the extract has a boiling point of less than the boiling point of the most volatile alcohol and this leaves the stripper as an overhead product and does not contaminate bottoms product, an important constituent of which is n-butanol. The n-butanol can be separated for use as a high-grade product. Thus, a particular advantage of this invention is that the separation results in a good product yield, not only of olefins and paraffins, but also of alcohols.

The process of the invention provides good separation between olefins, paraffins and oxygenates over a hydrocarbon material range of from C₄ to over C₂₀. The use of olefin-paraffin counter solvent avoids the complication of adding a second foreign solvent to the process. The boiling point of the counter-solvent is also important to ensure that n-butanol is separated from any counter-solvent in the extract stripper column. In addition, the process yields a substantially pure and useful alcohol product, as well as an olefin/paraffin product.

Example

A 3m vertical rotating disc extractor column, which is a multi-stage contacting device, was operated under conditions of liquid-liquid extraction. The column was fitted with feed points at the top bottom, and in-between. The solvent consisted of 18% water in acetonitrile. The solvent was fed at the top feed point at a feed rate of 2 kg/h. The counter-solvent was 1-octene (co-monomer

-9-

grade). The counter-solvent was introduced at the bottom feed point at a height of 2m at a rate of 1 kg/h. The feed was obtained from a condensate from a Fischer-Tropsch reaction in a slurry bed using a Fe/Mn/Zn catalyst operated at a pressure of 45 bar and a temperature of 230°C. The extraction operation was executed at 45°C. The solvent formed the continuous phase, and the interface was at the top of the column. For the particular set of accompanying results, the column's rotator was switched off.

Table 1 below shows the percent, by mass, of paraffins, olefins and alcohols in the hydrocarbon feed stream. The rest of the feed is made up from ketones, aldehydes, esters and isomers (branched alcohols, aldehydes, ketones, acids, esters and acetals).

Table 2 shows the percent, by mass, of paraffins, olefins and alcohols in the raffinate from the abovementioned extraction column. The rest of the composition is made up by ketones, aldehydes, esters and isomers.

Table 3 shows the percent, by mass, of alcohols in the extract. The rest of the extract is made up by aldehydes, ketones, esters and isomers.

The raffinate is then passed through a distillation column having a top temperature of 86°C, a bottom temperature of 200°C and 25 sieve trays. The distillation column is operated at a pressure of 150 kpa with a reflux ratio of 2. Table 4 below shows the composition of the overhead product and the bottoms product from this distillation column. It will be seen that the overhead product contains a high percentage of octene and octane which is recycled to the extraction column for use as the counter-solvent. The bottoms product is a mixture of olefins and paraffins, with low concentrations of oxygenates.

The extract is passed through a stripper having a top temperature of 88°C, a bottoms temperature of 106°C and 25 sieve trays. The stripper is operated at

-10-

a pressure of 150 kpa. The composition of the overhead product and the bottoms product from the stripper is shown in Table 4. It will be seen that the overhead product contains a high percentage of solvent (a mixture of acetonitrile and water), which is conveniently recycled to the extraction column. The bottoms product contains a high percentage of alcohols.

Table 1

Carbon No.	n-Paraffin Mass %	a-Olefins Mass %	1-Alcohols Mass %
3	0	0	0
4	0	0	1,2
5	0	0	4,2
6	0	0	4,2
7	0	0	3,9
8	0,3	0,3	3,2
9	3,8	3,3	2,3
10	4,3	3,8	1,5
11	4,0	3,4	0,9
12	3,4	2,7	0,6
13	2,7	2,3	0,3
14	2,1	1,8	0,1
15	1,6	1,2	0,1
16	1,1	0,7	0
17	0,8	0,4	0
18	0,5	0,3	0
19	0,3	0,1	0
20	0,2	0,1	0
21	0,1	0	0
22	0,1	0	0
Total	25,3	20,4	22,5

-11-

Table 2

Carbon No.	n-Paraffin Mass %	a-Olefins Mass %	1-Alcohols Mass %
6	0,1	0	0
7	0,1	0	0,2
8	*	*	0,4
9	5,1	3,4	0,3
10	10,0	6,4	0,5
11	9,4	5,4	0,6
12	8,4	4,1	0,5
13	6,8	2,8	0,4
14	5,1	1,7	0,3
15	3,5	0,9	0,2
16	2,3	0,5	0,2
17	1,5	0,2	0,1
18	1,0	0,1	0,1
19	0,7	0,1	0,1
20	0,5	0,1	0
21	0,3	0,1	0
22	0,3	0	0
23	0,2	0	0
24	0,2	0	0
25	0,1	0	0
26	0,1	0	0
27	0,1	0	0
28	0,1	0	0
29	0,1	0	0
Total	56,0	25,8	3,9

-12-

Table 3

Carbon No.	Alcohols Mass %
3	0,1
4	4,2
5	20,8
6	18,6
7	14,6
8	10,0
9	6,1
10	3,4
11	1,9
12	0,9
13	0,4
14	0,1
15	0,1
Total	81,2

Table 4

	Extract		Raffinate	
	Overhead	Bottoms	Overhead	Bottoms
MECN	72.2	0.0	6.7	0.0
WATER	17.0	0.0	0.6	0.0
BUOH	0.9	13.3	0.2	0.0
HEPTENE	0.2	0.0	2.1	0.0
NHEPTANE	0.4	0.0	4.4	0.0
PEOH	0.2	13.0		0.0
OCTENE	4.9	0.0	53.3	3.4
OCTANE	1.6	0.0	24.2	4.4
HEXOH	0.1	12.0	0.0	0.0
NONENE	0.1	0.0	0.2	4.2
NONANE	0.0		0.1	4.7
HEPOH	0.0	10.8	0.0	0.0
DECENE	0.0		0.0	3.0

-13-

DECANE	0.0		0.0	4.5
OCTOH	0.0	9.3	0.0	0.0
UNDECENE	0.0		0.0	3.2
UNDECANE	0.0		0.0	4.6
NONOH	0.0	8.4	0.0	0.0
DODECENE	0.0		0.0	2.6
DODECANE	0.0		0.0	4.4
DECANOL	0.0	7.0	0.0	0.0
TRIDECEN	0.0		0.0	2.5
TRIDECAN	0.0		0.0	4.7
UNDECOH	0.0	5.7	0.0	0.0
C14ENE	0.0		0.0	2.3
C14ANE	0.0		0.0	4.5
DODECOH	0.0	4.5	0.0	0.0
C15ENE	0.0		0.0	1.9
C15ANE	0.0		0.0	4.4
TRIDECOH	0.0	4.0	0.0	0.1
C16ENE	0.0		0.0	1.6
C16ANE	0.0		0.0	4.1
C14OH	0.0	3.0	0.0	0.4
C17ENE	0.0		0.0	1.3
C17ANE	0.0		0.0	3.9
C15OH	0.0	1.1	0.0	1.0
C18ENE	0.0		0.0	1.0
C18ANE	0.0		0.0	3.6
C16OH	0.0		0.0	1.2
C19ENE	0.0		0.0	0.8
C19ANE	0.0		0.0	3.3
C20ANE	0.0	0.0	0.0	3.5
C21ANE	0.0	0.0	0.0	3.0
C22ANE	0.0	0.0	0.0	2.4
C23ANE	0.0	0.0	0.0	2.0
C24ANE	0.0	0.0	0.0	1.5
C25ANE	0.0	0.0	0.0	1.2
C26ANE	0.0	0.0	0.0	0.8
C27ANE	0.0	0.0	0.0	0.6
C28ANE	0.0	0.0	0.0	0.4
C29ANE	0.0	0.0	0.0	0.3
C30ANE	0.0	0.0	0.0	0.3
Other	2.4	8.0	8.2	2.4
Total	100.0	100.0	100.0	100.0

CLAIMS

1. A method for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream including hydrocarbons and oxygenates, the method including contacting the hydrocarbon stream under conditions of liquid-liquid extraction with a polar solvent and a non-polar organic counter-solvent, wherein the organic counter-solvent comprises olefins and/or paraffins.
2. The method according to claim 1 wherein the organic counter-solvent is selected to have a boiling point which is less than the boiling point of the most volatile alcohol in the hydrocarbon stream.
3. The method according to claim 1 or 2 wherein the hydrocarbon stream is passed through a distillation column prior to introduction to the liquid-liquid extraction, to remove C₁ to C₇ hydrocarbons and C₁ to C₃ alcohols from the stream.
4. The method according to claim 3 wherein the organic counter-solvent comprises olefins and/or paraffins in the C₈ range.
5. The method according to any one of claims 3 to 4 wherein the C₂₁ – plus hydrocarbons are removed from the hydrocarbon stream by distillation prior to the liquid-liquid extraction so that the stream comprises C₄ to C₂₀ hydrocarbons.
6. The method according to claim 5 wherein the C₁₉ – plus hydrocarbons are removed from the hydrocarbon stream by distillation prior to liquid-liquid extraction so that the stream comprises C₄ to C₁₈ hydrocarbons.

-15-

7. The method according to claim 5 or 6 wherein the polar solvent is a light polar solvent.
8. The method according to claim 7 wherein the solvent comprises a mixture of water and an organic liquid.
9. The method according to claim 8 wherein the light polar solvent is a mixture of water and acetonitrile.
10. The method according to claim 9 wherein the water comprises no more than 19% of the solvent.
11. The method according to any one of the preceding claims wherein a raffinate from the liquid-liquid extractor is introduced into a distillation column and a mixture of paraffins and olefins is recovered as a bottoms product from the distillation column.
12. The method of any one of the preceding claims wherein the overhead product from the distillation column is recycled to the lower counter-solvent inlet of the extraction column.
13. The method according to any one of the preceding claims wherein an extract from the liquid-liquid extractor is sent to a stripping column, and an overhead product from the stripping column is recycled to the upper solvent inlet of the extraction column.
14. The method according to any one of the preceding claims wherein the bottoms product from the stripping column is an oxygenate stream containing low concentrations of non-polar material.

-16-

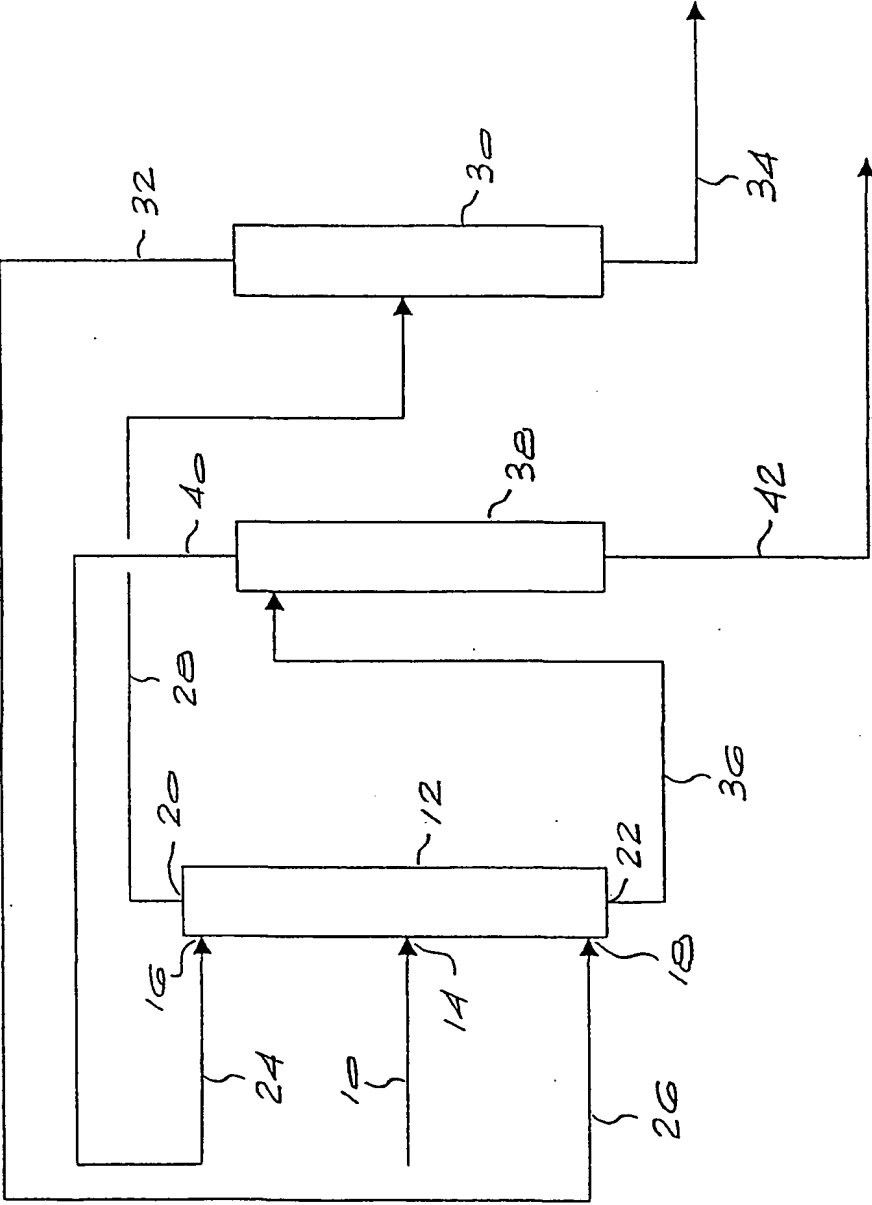
15. A method for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream comprising C_4 to over C_{20} hydrocarbons and oxygenates, the method including the steps of:

introducing the hydrocarbon stream into a vertical liquid-liquid extraction column, the liquid extraction column including an upper solvent inlet, a lower counter-solvent inlet, an upper raffinate outlet and lower extract outlet;

introducing a polar solvent into the extraction column via the upper solvent inlet; and

introducing a non-polar counter-solvent into the extraction column via the lower counter-solvent inlet.

16. The method of claim 15 wherein the hydrocarbon stream is added to the liquid-liquid extraction column, along the liquid extraction column.



(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
18 April 2002 (18.04.2002)

PCT

(10) International Publication Number
WO 02/031085 A3

(51) International Patent Classification⁷: **C10G 21/04**

(74) Agents: **GILSON, David, Grant et al.**; Spoor and Fisher,
P.O. Box 41312, 2024 Craighall (ZA).

(21) International Application Number: PCT/IB01/01874

(22) International Filing Date: 9 October 2001 (09.10.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
2000/5508 9 October 2000 (09.10.2000) ZA
60/239,060 9 October 2000 (09.10.2000) US

(71) Applicant (for all designated States except US): **SASOL TECHNOLOGY (PTY) LTD.** [ZA/ZA]; 1 Klasie Havenga Road, 1947 Sasolburg (ZA).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(72) Inventors; and

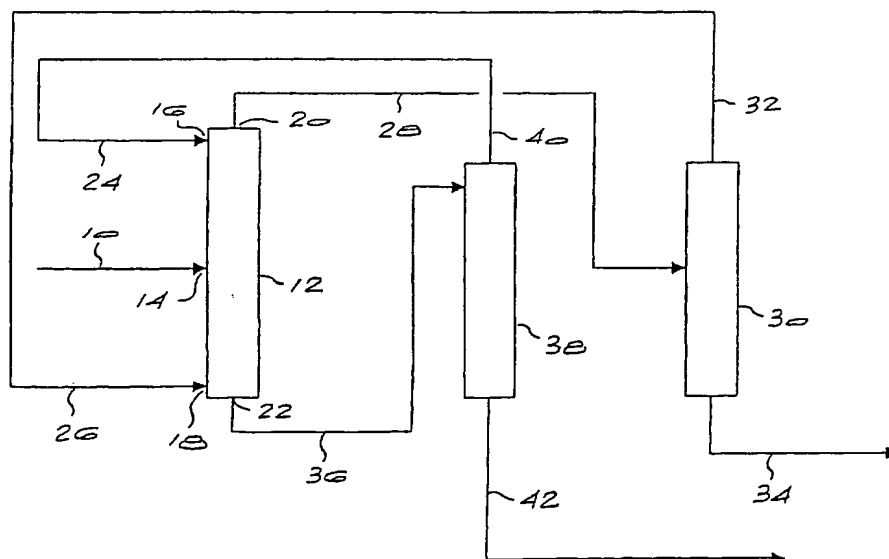
(75) Inventors/Applicants (for US only): **DE WET, Johan, Peter** [ZA/ZA]; 65 Beethoven Street, 1911 Vanderbijlpark (ZA). **SCHOLTZ, Jacob, Johannes** [ZA/ZA]; 6 Oosthuizen Street, 1947 Sasolburg (ZA).

Published:

— with international search report

[Continued on next page]

(54) Title: SEPARATION OF OXYGENATES FROM A HYDROCARBON STREAM



(57) Abstract: This invention relates to a method for separating olefins and paraffins from oxygenates in a liquid hydrocarbon stream containing a high proportion of olefins, paraffins and oxygenates (mainly alcohols). Typically, the hydrocarbon stream is obtained from a Fischer-Tropsch process. The organic counter-solvent has a boiling point which is less than the boiling point of the most volatile alcohol in the hydrocarbon stream. A raffinate from the liquid-liquid extractor is passed to a distillation column. A bottoms product from the distillation column comprises olefins and paraffins, and the overhead product comprising solvents is recycled. An extract from the liquid-liquid extractor is sent to a stripping column, where a bottoms product containing pure alcohol is obtained. The overhead product containing counter-solvent is recycled.

WO 02/031085 A3



— *before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments*

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(88) Date of publication of the international search report:
7 November 2002

INTERNATIONAL SEARCH REPORT

International Application No

PCT/IB 01/01874

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C10G21/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C10G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 717 725 A (UNIVERSAL OIL PROD CO) 3 November 1954 (1954-11-03) page 1, line 13 -page 4, line 26	1-8, 12, 15, 16
Y	page 9, line 100 - line 114; claim 1; figure 1 page 8, line 111 - line 127	1-14
X	GB 661 916 A (BATAAFSCHE PETROLEUM) 28 November 1951 (1951-11-28) page 1, line 65 - line 76; claim 1; example 1	1, 15, 16
Y		1, 11-14
Y	WO 99 58625 A (VILLIERS WALDO EUGENE DE ;NAUDE HUBERT (ZA); WET PETRA DE (ZA); HO) 18 November 1999 (1999-11-18) claims 1,3,4,7,13; figure 1	1-10, 13, 14
	--- -/--	



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *G* document member of the same patent family

Date of the actual completion of the international search

26 August 2002

Date of mailing of the international search report

30/08/2002

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Deurinck, P

INTERNATIONAL SEARCH REPORT

International Application No

PCT/IB 01/01874

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 3 725 255 A (BARILLI F ET AL) 3 April 1973 (1973-04-03) figure 1 -----	1,11,12

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/IB 01/01874

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
GB 717725	A	03-11-1954	NONE	
GB 661916	A	28-11-1951	NONE	
WO 9958625	A	18-11-1999	AU 3438899 A BR 9910299 A CN 1300315 T EP 1092005 A1 WO 9958625 A1 JP 2002514680 T	29-11-1999 25-09-2001 20-06-2001 18-04-2001 18-11-1999 21-05-2002
US 3725255	A	03-04-1973	DE 2123292 A1 FR 2088511 A1 GB 1308735 A JP 50008722 B NL 7106509 A	25-11-1971 07-01-1972 07-03-1973 07-04-1975 16-11-1971